

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF

ROBL ET AL.

APPLICATION NO: NOT ASSIGNED YET

FILED: HERewith

FOR: METHOD FOR TREATING ATHEROSCLEROSIS EMPLOYING  
AN AP2 INHIBITOR AND COMBINATION

Assistant Commissioner for Patents  
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

Kindly amend the above-referenced application as follows:

In the Specification:

Page 1, line 3, please insert the following:

--This application is a continuation of U.S. Serial No. 09/390,275 filed September 7, 1999, which claims priority from U.S. Provisional Application Serial No. 60/100,677. The entirety of each of these applications is incorporated herein by reference.--

Page 3, please replace the paragraph beginning at line 1 and ending at line 16 with the following:

--The aP2 inhibitors suitable for use in the method of the invention are compounds which bind to the aP2 protein (SEQ ID NO:1) and inhibits its function and/or its ability to bind free fatty acids. The compounds will preferably contain less than 60 carbon atoms, more preferably less than 45 carbon atoms, and will contain less than 20 heteroatoms, more preferably less than 12 heteroatoms. They contain a hydrogen bond donator or acceptor group, preferably acidic in nature, which includes, but is not limited to, CO<sub>2</sub>H, tetrazole, SO<sub>3</sub>H, PO<sub>3</sub>H, P(R)(O)OH (where R is lower

alkyl or lower alkoxy), OH, NHSO<sub>2</sub>R' or CONHSO<sub>2</sub>R' (where R' is lower alkyl), and thiazolidindione, and interacts (directly or through an intervening water molecule), either by ionic or hydrogen bonding interactions, with one, two, or three of the three amino acid residues, designated as Arg 106, Arg 126 and Tyr 128 in human aP2, within the aP2 protein (SEQ ID NO:1).--

Page 3, please replace the paragraph beginning at line 25 and ending at line 32 with the following:

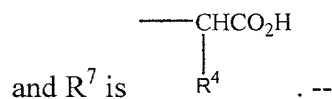
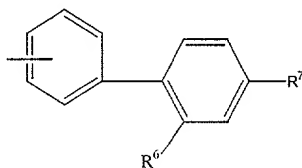
--The hydrophobic substituent binds to (in) and/or interacts with a discrete pocket within the aP2 protein (SEQ ID NO:1) defined roughly by the amino acid residues Phe 16, Tyr 19, Met 20, Val 23, Val 25, Ala 33, Phe 57, Thr 74, Ala 75, Asp 76, Arg 78 in human aP2. The through space distance from the hydrogen bond donor/acceptor group and the additional substituent group is within the distance of about 7 to about 15 Angstroms.--

Page 4, please replace the paragraph beginning at line 11 and ending at line 13 with the following:

-- The accompanying Figure 1 is a computer generated image of a partial X-ray structure of compound XVIA (described hereinafter) bound to human aP2.--

Page 6, please replace the paragraph beginning at line 22 and ending at line 26 with the following:

--Preferred are the examples where A is defined as above and B is



Page 28, please replace the paragraph beginning at line 21 and ending at line 29 with the following:

--The other antiatherosclerotic agent employed in the methods of the invention include MTP inhibitors disclosed in U.S. Patent Nos. 5,595,872, 5,739,135, 5,712,279, 5,760,246, 5,827,875, 5,885,983 and 5,962,440. Preferred are each of the preferred MTP inhibitors disclosed in each of the above patents.--

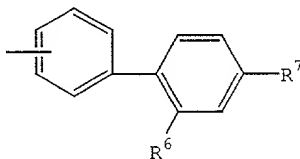
In the Claims:

3. (Amended) The method as defined in Claim 1 wherein the aP2 inhibitor contains a hydrogen bond donator or acceptor group and interacts directly or through an intervening water molecule either by ionic or hydrogen bonding interactions, with one, two, or three of the three amino acid residues, designated as Arg 106, Arg 126 and Tyr 128 in human aP2 within the aP2 protein (SEQ ID NO:1).

Attached hereto is a marked-up version of the changes made to the claims by the current amendment. The attached page is captioned "**Version With Marking to Show Changes Made**". These amendments add no new matter.

REMARKS

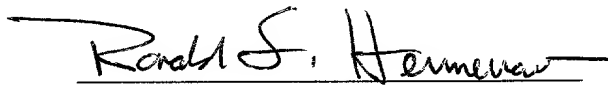
Applicants have amended the paragraph beginning at page 6, line 22 of the specification to insert the following chemical structure:



Support for this amendment can be found in the priority document (U.S. provisional application Serial No. 60/100,677, filed September 17, 1998) at page 5 line 24.

Respectfully submitted,

Bristol-Myers Squibb Company  
Patent Department  
P.O. Box 4000  
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Date: 7/13/01

  
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Attorney for Applicants  
Reg. No. 34,620

Version with Markings to Show Changes Made

Page 1, line 3, has been amended as follows please see attachment for marked up version.

This application is a continuation of U.S. Serial No. 09/390,275 filed September 7, 1999, which claims priority from U.S. Provisional Application Serial No. 60/100,677. The entirety of each of these applications is incorporated herein by reference.

Page 3, the paragraph beginning at line 1 and ending at line 16 has been amended as follows:

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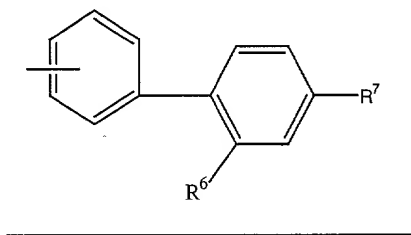
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October 20, 1998, now U.S. Patent No.] 5,962,440. Preferred are each of the preferred MTP inhibitors disclosed in each of the above patents[ and applications].

In the Claims:

3. (Amend) The method as defined in Claim 1 wherein the aP2 inhibitor contains a hydrogen bond donator or acceptor group and interacts directly or through an intervening water molecule either by ionic or hydrogen bonding interactions, with one, two, or three of the three amino acid residues, designated as Arg 106, Arg 126 and Tyr 128 in human aP2 within the aP2 protein (SEQ ID NO:1).

This application is a continuation of U.S. Serial No. 09/390,275 filed September 7, 1999, which claims priority from U.S. Provisional Application Serial No. 60/100,677. The entirety of each of these applications is incorporated herein by reference.

METHOD FOR TREATING ATHEROSCLEROSIS EMPLOYING

AN aP2 INHIBITOR AND COMBINATION

Field of the Invention

- 5 The present invention relates to a method for treating atherosclerosis and related diseases, employing an aP2 inhibitor alone or in combination with another type antiatherosclerotic agent.

Background of the Invention

- 10 Fatty acid binding proteins (FABPs) are small cytoplasmic proteins which bind to fatty acids such as oleic acids which are important metabolic fuels and cellular regulators. Dysregulation of fatty acid  
15 metabolism in adipose tissue is a prominent feature of insulin resistance and the transition from obesity to non-insulin dependent diabetes mellitus (NIDDM or Type II diabetes).

- aP2, an abundant 14.6 KDa cytosolic protein in  
20 adipocytes, and one of a family of homologous intracellular fatty acid binding proteins (FABPs), is involved in the regulation of fatty acid trafficking in adipocytes and mediates fatty acid fluxes in adipose tissue. G.S. Hotamisligil et al, "Uncoupling of Obesity from Insulin  
25 Resistance Through a Targeted Mutation in aP2, the Adipocyte Fatty Acid Binding Protein", Science, Vol. 274, Nov. 22, 1996, pp. 1377-1379, report that aP2-deficient mice placed on a high fat diet for several weeks developed dietary obesity, but, unlike control-mice on a similar  
30 diet, did not develop insulin resistance or diabetes. Hotamisligil et al conclude that "aP2 is central to the pathway that links obesity to insulin resistance" (Abstract, page 1377).

- DIALOG ALERT DBDR928 dates January 2, 1997,  
35 Pharmaprojects No. 5149 (Knight-Ridder Information) discloses that a major drug company "is using virtual screening techniques to identify potential new antidiabetic